1: NP_000168. Reports gelsolin isoform ...[gi:4504165]

Comment Features Sequence

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LOCUS       NP_000168                782 aa            linear   PRI 31-OCT-2000
DEFINITION  gelsolin (amyloidosis, Finnish type); Gelsolin [Homo sapiens].
ACCESSION   NP_000168
VERSION     NP_000168.1  GI:4504165
DBSOURCE    REFSEQ: accession NM_000177.1
KEYWORDS    .
SOURCE      Homo sapiens (human)
  ORGANISM  Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1  (residues 1 to 782)
  AUTHORS   Hiltunen T, Kiuru S, Hongell V, Helio T, Palo J and Peltonen L.
  TITLE     Finnish type of familial amyloidosis: cosegregation of
            Asp187----Asn mutation of gelsolin with the disease in three large
            families
  JOURNAL   Am. J. Hum. Genet. 49 (3), 522-528 (1991)
   PUBMED   1652889
REFERENCE   2  (residues 1 to 782)
  AUTHORS   Kwiatkowski,D.J., Stossel,T.P., Orkin,S.H., Mole,J.E., Colten,H.R.
            and Yin,H.L.
  TITLE     Plasma and cytoplasmic gelsolins are encoded by a single gene and
            contain a duplicated actin-binding domain
  JOURNAL   Nature 323 (6087), 455-458 (1986)
   PUBMED   3020431
COMMENT     PROVISIONAL REFSEQ: This record has not yet been subject to final
            NCBI review. The reference sequence was derived from X04412.1.
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